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|------|----|--------|--|
| NEWS | 1 | | Web Page URLs for STN Seminar Schedule - N. America |
| NEWS | 2 | Apr 08 | "Ask CAS" for self-help around the clock |
| NEWS | 3 | Apr 09 | BEILSTEIN: Reload and Implementation of a New Subject Area |
| NEWS | 4 | Apr 09 | ZDB will be removed from STN |
| NEWS | 5 | Apr 19 | US Patent Applications available in IFICDB, IFIPAT, and IFIUDB |
| NEWS | 6 | Apr 22 | Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS |
| NEWS | 7 | Apr 22 | BIOSIS Gene Names now available in TOXCENTER |
| NEWS | 8 | Apr 22 | Federal Research in Progress (FEDRIP) now available |
| NEWS | 9 | Jun 03 | New e-mail delivery for search results now available |
| NEWS | 10 | Jun 10 | MEDLINE Reload |
| NEWS | 11 | Jun 10 | PCTFULL has been reloaded |
| NEWS | 12 | Jul 02 | FOREGE no longer contains STANDARDS file segment |
| NEWS | 13 | Jul 22 | USAN to be reloaded July 28, 2002; saved answer sets no longer valid |
| NEWS | 14 | Jul 29 | Enhanced polymer searching in REGISTRY |
| NEWS | 15 | Jul 30 | NETFIRST to be removed from STN |
| NEWS | 16 | Aug 08 | CANCERLIT reload |
| NEWS | 17 | Aug 08 | PHARMAMarketLetter(PHARMAML) - new on STN |
| NEWS | 18 | Aug 08 | NTIS has been reloaded and enhanced |
| NEWS | 19 | Aug 19 | Aquatic Toxicity Information Retrieval (AQUIRE) now available on STN |
| NEWS | 20 | Aug 19 | IFIPAT, IFICDB, and IFIUDB have been reloaded |
| NEWS | 21 | Aug 19 | The MEDLINE file segment of TOXCENTER has been reloaded |
| NEWS | 22 | Aug 26 | Sequence searching in REGISTRY enhanced |
| NEWS | 23 | Sep 03 | JAPIO has been reloaded and enhanced |
| NEWS | 24 | Sep 16 | Experimental properties added to the REGISTRY file |
| NEWS | 25 | Sep 16 | CA Section Thesaurus available in CAPLUS and CA |
| NEWS | 26 | Oct 01 | CASREACT Enriched with Reactions from 1907 to 1985 |
| NEWS | 27 | Oct 21 | EVENTLINE has been reloaded |
| NEWS | 28 | Oct 24 | BEILSTEIN adds new search fields |
| NEWS | 29 | Oct 24 | Nutraceuticals International (NUTRACEUT) now available on STN |
| NEWS | 30 | Oct 25 | MEDLINE SDI run of October 8, 2002 |
| NEWS | 31 | Nov 18 | DKILIT has been renamed APOLLIT |
| NEWS | 32 | Nov 25 | More calculated properties added to REGISTRY |
| NEWS | 33 | Dec 02 | TIBKAT will be removed from STN |
| NEWS | 34 | Dec 04 | CSA files on STN |
| NEWS | 35 | Dec 17 | PCTFULL now covers WP/PCT Applications from 1978 to date |
| NEWS | 36 | Dec 17 | TOXCENTER enhanced with additional content |
| NEWS | 37 | Dec 17 | Adis Clinical Trials Insight now available on STN |
| NEWS | 38 | Dec 30 | ISMEC no longer available |
| NEWS | 39 | Jan 13 | Indexing added to some pre-1967 records in CA/CAPLUS |
| NEWS | 40 | Jan 21 | NUTRACEUT offering one free connect hour in February 2003 |
| NEWS | 41 | Jan 21 | PHARMAML offering one free connect hour in February 2003 |
| NEWS | 42 | Jan 29 | Simultaneous left and right truncation added to COMPENDEX, ENERGY, INSPEC |
| NEWS | 43 | Feb 13 | CANCERLIT is no longer being updated |
| NEWS | 44 | Feb 24 | METADEX enhancements |
| NEWS | 45 | Feb 24 | PCTGEN now available on STN |

NEWS 46 Feb 24 TEMA now available on STN
 NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation
 NEWS 48 Feb 26 PCTFULL now contains images
 NEWS 49 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results

NEWS EXPRESS January 6 CURRENT WINDOWS VERSION IS V6.01a,
 CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
 AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002
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 NEWS INTER General Internet Information
 NEWS LOGIN Welcome Banner and News Items
 NEWS PHONE Direct Dial and Telecommunication Network Access to STN
 NEWS WWW CAS World Wide Web Site (general information)

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FILE 'MEDLINE' ENTERED AT 15:07:48 ON 14 MAR 2003

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=> s integrative transformants

L1 41 INTEGRATIVE TRANSFORMANTS

=> s reiterated ribosomal DNA
L2 0 REINTERATED RIBOSOMAL DNA

=> s ribosomal DNA
L3 16927 RIBOSOMAL DNA

=> s l3 and reiterated
L4 50 L3 AND REITERATED

=> s l1 and l4
L5 0 L1 AND L4

=> s l4 and yeast
L6 13 L4 AND YEAST

=> d l6 ti abs ibib tot

L6 ANSWER 1 OF 13 MEDLINE

TI Replicon size of **yeast ribosomal DNA**.

AB The ribosomal RNAs of the **yeast** *Saccharomyces cerevisiae* are transcribed from a 9K bp stretch of DNA which is **reiterated** about 120-fold in a continuous array, about 360 microns long, on chromosome XII. Although ARS activity has been detected in the repeat unit, the size and disposition of replicons along this array of identical genes has not hitherto been determined. We have used immobilised rRNA as a probe to examine the size of radioactively labelled rDNA replicons resolved on alkaline sucrose gradients. The replicons were found to be uniformly sized, about 5 repeat units in length, and groups of 4 adjacent replicons may be activated simultaneously. These observations suggest that replicon initiation events are not determined solely by the recognition of specific DNA sequences that function as origins of replication.

ACCESSION NUMBER: 85035837 MEDLINE

DOCUMENT NUMBER: 85035837 PubMed ID: 6387390

TITLE: Replicon size of **yeast ribosomal DNA**.

AUTHOR: Walmsley R M; Johnston L H; Williamson D H; Oliver S G

SOURCE: MOLECULAR AND GENERAL GENETICS, (1984) 195 (1-2) 260-6.

Journal code: 0125036. ISSN: 0026-8925.

PUB. COUNTRY: GERMANY, WEST: Germany, Federal Republic of

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198411

ENTRY DATE: Entered STN: 19900320

Last Updated on STN: 19900320

Entered Medline: 19841126

L6 ANSWER 2 OF 13 MEDLINE

TI Simple Mendelian inheritance of the **reiterated ribosomal DNA of yeast**.

AB A diploid strain of **yeast** (*Saccharomyces cerevisiae*) was found to be heterozygous for two forms of the highly repetitious **ribosomal DNA**. These forms could be distinguished by the pattern of fragments produced after digestion with the site-specific restriction endonuclease EcoRI. The mode of inheritance of **ribosomal DNA** was determined by tetrad analysis. Of 14 tetrads analyzed, 12 clearly showed the **ribosomal DNA** forms segregating as a single Mendelian unit. The simplest interpretation of this result is that all of the approximately 100 copies of the **ribosomal DNA** genes of the **yeast** cell are located on one chromosome and that meiotic recombination within these genes is suppressed. Two of the 14 tetrads showed the segregation patterns expected as the result of mitotic recombination within the **ribosomal DNA**.

ACCESSION NUMBER: 78053057 MEDLINE
 DOCUMENT NUMBER: 78053057 PubMed ID: 337310
 TITLE: Simple Mendelian inheritance of the **reiterated ribosomal DNA** of yeast.
 AUTHOR: Petes T D; Botstein D
 SOURCE: PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA, (1977 Nov) 74 (11) 5091-5.
 Journal code: 7505876. ISSN: 0027-8424.
 PUB. COUNTRY: United States
 DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)
 LANGUAGE: English
 FILE SEGMENT: Priority Journals
 ENTRY MONTH: 197801
 ENTRY DATE: Entered STN: 19900314
 Last Updated on STN: 19900314
 Entered Medline: 19780127

L6 ANSWER 3 OF 13 WPIDS (C) 2003 THOMSON DERWENT
 TI **Yeast** which ferments xylose to ethanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites.

AN 1997-558974 [51] WPIDS

AB WO 9742307 A UPAB: 19991020

Novel **yeast** which ferments xylose to ethanol, comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA** sites; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol, where the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations.

USE - The methods can produce **yeast**, which even upon culture in non-selective medium for multiple generations, e.g. up to 20, retain their full capability to ferment xylose to ethanol.

Dwg.0/12

ACCESSION NUMBER: 1997-558974 [51] WPIDS

DOC. NO. CPI: C1997-178545

TITLE: **Yeast** which ferments xylose to ethanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites.

DERWENT CLASS: D16 D17 E17 H06

INVENTOR(S): CHEN, Z; HO, N W Y

PATENT ASSIGNEE(S): (PURD) PURDUE RES FOUND

COUNTRY COUNT: 76

PATENT INFORMATION:

| PATENT NO | KIND | DATE | WEEK | LA | PG |
|---|------|----------|-----------|----|----|
| WO 9742307 | A1 | 19971113 | (199751)* | EN | 66 |
| RW: AT BE CH DE DK EA ES FI FR GB GH GR IE IT KE LS LU MC MW NL OA PT SD SE SZ UG | | | | | |
| W: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB GE HU IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK TJ TM TR TT UA UG US UZ VN YU | | | | | |
| AU 9728301 | A | 19971126 | (199813) | | |
| EP 898616 | A1 | 19990303 | (199913) | EN | |
| R: AT BE DE DK ES FI FR GB GR IE IT NL PT SE | | | | | |
| CN 1225125 | A | 19990804 | (199949) | | |

JP 2000509988 W 20000808 (200043) 50
 MX 9809223 A1 19990701 (200061)
 AU 731102 B 20010322 (200122)
 BR 9710963 A 20010731 (200146)

APPLICATION DETAILS:

| PATENT NO | KIND | APPLICATION | DATE |
|---------------|------|----------------|----------|
| WO 9742307 | A1 | WO 1997-US7663 | 19970506 |
| AU 9728301 | A | AU 1997-28301 | 19970506 |
| EP 898616 | A1 | EP 1997-922698 | 19970506 |
| | | WO 1997-US7663 | 19970506 |
| CN 1225125 | A | CN 1997-196195 | 19970506 |
| JP 2000509988 | W | JP 1997-540153 | 19970506 |
| | | WO 1997-US7663 | 19970506 |
| MX 9809223 | A1 | MX 1998-9223 | 19981105 |
| AU 731102 | B | AU 1997-28301 | 19970506 |
| BR 9710963 | A | BR 1997-10963 | 19970506 |
| | | WO 1997-US7663 | 19970506 |

FILING DETAILS:

| PATENT NO | KIND | PATENT NO |
|---------------|------------------|------------|
| AU 9728301 | A Based on | WO 9742307 |
| EP 898616 | A1 Based on | WO 9742307 |
| JP 2000509988 | W Based on | WO 9742307 |
| AU 731102 | B Previous Publ. | AU 9728301 |
| | Based on | WO 9742307 |
| BR 9710963 | A Based on | WO 9742307 |

PRIORITY APPLN. INFO: US 1996-16865P 19960506

L6 ANSWER 4 OF 13 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.

TI Simple Mendelian inheritance of the **reiterated ribosomal DNA of yeast.**

ACCESSION NUMBER: 78243218 EMBASE

DOCUMENT NUMBER: 1978243218

TITLE: Simple Mendelian inheritance of the **reiterated ribosomal DNA of yeast.**

AUTHOR: Petes T.D.; Botstein D.

CORPORATE SOURCE: Dept. Biol., MIT, Cambridge, Mass. 02139, United States

SOURCE: Proceedings of the National Academy of Sciences of the United States of America, (1977) 74/11 (5091-5095).

CODEN: PNASA6

COUNTRY: United States

DOCUMENT TYPE: Journal

FILE SEGMENT: 022 Human Genetics

LANGUAGE: English

L6 ANSWER 5 OF 13 DGENE (C) 2003 THOMSON DERWENT

TI **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

AN AAV12824 DNA DGENE

AB This sequence represents an amplification primer for the **yeast** 5S rDNA sequence. The amplified sequence can be used in the **yeast** of the invention, which ferments xylose to ethanol. The **yeast** comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA** sites; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to

non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol; the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations. The **yeast** is produced by integrating multiple copies of exogenous DNA into **reiterated** chromosomal DNA of cells. The **yeast** produced by the integration method, even upon culture in non-selective medium for multiple generations (e.g. up to 20), retain their full capability to ferment xylose to ethanol.

ACCESSION NUMBER: AAV12824 DNA DGENE

TITLE: **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

INVENTOR: Chen Z; Ho N W Y

PATENT ASSIGNEE: (PURD)PURDUE RES FOUND.

PATENT INFO: WO 9742307 A1 19971113 66p

APPLICATION INFO: WO 1997-US7663 19970506

PRIORITY INFO: US 1996-16865 19960506

DOCUMENT TYPE: Patent

LANGUAGE: English

OTHER SOURCE: 1997-558974 [51]

L6 ANSWER 6 OF 13 DGENE (C) 2003 THOMSON DERWENT

TI **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

AN AAV12829 DNA DGENE

AB This sequence is an amplification primer for the **yeast** Tn903 kanamycin resistance gene. The amplified sequence can be used in the **yeast** of the invention, which ferments xylose to ethanol. The **yeast** comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA** sites; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol; the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations. The **yeast** is produced by integrating multiple copies of exogenous DNA into **reiterated** chromosomal DNA of cells. The **yeast** produced by the integration method, even upon culture in non-selective medium for multiple generations (e.g. up to 20), retain their full capability to ferment xylose to ethanol.

ACCESSION NUMBER: AAV12829 DNA DGENE

TITLE: **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

INVENTOR: Chen Z; Ho N W Y

PATENT ASSIGNEE: (PURD)PURDUE RES FOUND.

PATENT INFO: WO 9742307 A1 19971113 66p

APPLICATION INFO: WO 1997-US7663 19970506

PRIORITY INFO: US 1996-16865 19960506

DOCUMENT TYPE: Patent

LANGUAGE: English

OTHER SOURCE: 1997-558974 [51]

L6 ANSWER 7 OF 13 DGENE (C) 2003 THOMSON DERWENT

TI **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

AN AAV12828 DNA DGENE

AB This sequence is an amplification primer for the **yeast** Tn903 kanamycin resistance gene. The amplified sequence can be used in the **yeast** of the invention, which ferments xylose to ethanol. The **yeast** comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA** sites; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol; the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations. The **yeast** is produced by integrating multiple copies of exogenous DNA into **reiterated** chromosomal DNA of cells. The **yeast** produced by the integration method, even upon culture in non-selective medium for multiple generations (e.g. up to 20), retain their full capability to ferment xylose to ethanol.

ACCESSION NUMBER: AAV12828 DNA DGENE

TITLE: **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

INVENTOR: Chen Z; Ho N W Y

PATENT ASSIGNEE: (PURD) PURDUE RES FOUND.

PATENT INFO: WO 9742307 A1 19971113 66p

APPLICATION INFO: WO 1997-US7663 19970506

PRIORITY INFO: US 1996-16865 19960506

DOCUMENT TYPE: Patent

LANGUAGE: English

OTHER SOURCE: 1997-558974 [51]

L6 ANSWER 8 OF 13 DGENE (C) 2003 THOMSON DERWENT

TI **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA** sites

AN AAV12827 DNA DGENE

AB This sequence is an amplification primer for the **yeast** Tn903 kanamycin resistance gene. The amplified sequence can be used in the **yeast** of the invention, which ferments xylose to ethanol. The **yeast** comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA** sites; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol; the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations. The **yeast** is produced by integrating multiple copies of exogenous DNA into **reiterated** chromosomal DNA of cells. The **yeast** produced by the integration method, even upon culture in non-selective medium for multiple generations (e.g. up to 20), retain their full capability to ferment xylose to ethanol.

ACCESSION NUMBER: AAV12827 DNA DGENE

TITLE: **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and

xylulokinase genes integrated at each of its multiple
reiterated ribosomal DNA sites
INVENTOR: Chen Z; Ho N W Y
PATENT ASSIGNEE: (PURD)PURDUE RES FOUND.
PATENT INFO: WO 9742307 A1 19971113 66p
APPLICATION INFO: WO 1997-US7663 19970506
PRIORITY INFO: US 1996-16865 19960506
DOCUMENT TYPE: Patent
LANGUAGE: English
OTHER SOURCE: 1997-558974 [51]

L6 ANSWER 9 OF 13 DGENE (C) 2003 THOMSON DERWENT

TI **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA sites**

AN AAV12826 DNA DGENE

AB This sequence is an amplification primer for the **yeast** Tn903 kanamycin resistance gene. The amplified sequence can be used in the **yeast** of the invention, which ferments xylose to ethanol. The **yeast** comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA sites**; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol; the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations. The **yeast** is produced by integrating multiple copies of exogenous DNA into **reiterated** chromosomal DNA of cells. The **yeast** produced by the integration method, even upon culture in non-selective medium for multiple generations (e.g. up to 20), retain their full capability to ferment xylose to ethanol.

ACCESSION NUMBER: AAV12826 DNA DGENE

TITLE: **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA sites**

INVENTOR: Chen Z; Ho N W Y

PATENT ASSIGNEE: (PURD)PURDUE RES FOUND.

PATENT INFO: WO 9742307 A1 19971113 66p

APPLICATION INFO: WO 1997-US7663 19970506

PRIORITY INFO: US 1996-16865 19960506

DOCUMENT TYPE: Patent

LANGUAGE: English

OTHER SOURCE: 1997-558974 [51]

L6 ANSWER 10 OF 13 DGENE (C) 2003 THOMSON DERWENT

TI **Yeast** which ferments xylose to methanol - comprising xylitol reductase, xylitol dehydrogenase and xylulokinase genes integrated at each of its multiple **reiterated ribosomal DNA sites**

AN AAV12825 DNA DGENE

AB This sequence represents an amplification primer for the **yeast** 5S rDNA sequence. The amplified sequence can be used in the **yeast** of the invention, which ferments xylose to ethanol. The **yeast** comprises: (a) xylose reductase (XR), xylitol dehydrogenase (XD) and xylulokinase (XK) genes integrated at each of its multiple **reiterated ribosomal DNA sites**; (b) multiple copies of exogenous DNA, including XR, XD, and XK genes, fused to non-glucose inhibited promoters integrated into its chromosomal DNA, where the **yeast** simultaneously ferments glucose and xylose to

ethanol; or (c) multiple copies of an introduced DNA containing XR, XD and XK genes, where the **yeast** ferments xylose to ethanol; the yeasts of (b) and (c) retain their capacity for fermenting xylose to ethanol when cultured under non-selective conditions for at least 20 generations. The **yeast** is produced by integrating multiple copies of exogenous DNA into **reiterated** chromosomal DNA of cells. The **yeast** produced by the integration method, even upon culture in non-selective medium for multiple generations (e.g. up to 20), retain their full capability to ferment xylose to ethanol.

ACCESSION NUMBER: AAV12825 DNA DGENE
TITLE: **Yeast** which ferments xylose to methanol -
comprising xylitol reductase, xylitol dehydrogenase and
xylulokinase genes integrated at each of its multiple
reiterated ribosomal DNA sites
INVENTOR: Chen Z; Ho N W Y
PATENT ASSIGNEE: (PURD)PURDUE RES FOUND.
PATENT INFO: WO 9742307 A1 19971113 66p
APPLICATION INFO: WO 1997-US7663 19970506
PRIORITY INFO: US 1996-16865 19960506
DOCUMENT TYPE: Patent
LANGUAGE: English
OTHER SOURCE: 1997-558974 [51]

L6 ANSWER 11 OF 13 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
TI THE MAJOR PROMOTER ELEMENT OF RIBOSOMAL RNA TRANSCRIPTION IN **YEAST**
LIES 2 KILOBASE PAIR UPSTREAM.
AB Conventional genetic analysis of the transcription of rDNA [**ribosomal DNA**] in **yeast** is precluded because the genes are highly **reiterated**. As an alternative strategy to determine which sequences modulate transcription of pre-rRNA, a series of artificial rRNA genes containing a fragment of DNA from Escherichia coli bacteriophage T7 were introduced into the **yeast** Saccharomyces cerevisiae. Correct transcription of the artificial genes was observed. Three regions of ribosomal spacer affected transcription of rRNA. Sequences within 210 bp [base pair] of the 5' terminus of 35S rRNA support low levels of transcription, but at multiple initiation points. Sequences from -210 to -2230 direct correct initiation and increase somewhat the efficiency of transcription. Most striking is that sequences from -2230 to -2420 stimulate transcription 15-fold. The function of this major promoter element is absolutely orientation-dependent but relatively independent of position. Its activity is blocked when an rRNA transcription termination sequence is placed between it and the site of initiation.

ACCESSION NUMBER: 1985:278527 BIOSIS
DOCUMENT NUMBER: BA79:58523
TITLE: THE MAJOR PROMOTER ELEMENT OF RIBOSOMAL RNA TRANSCRIPTION
IN **YEAST** LIES 2 KILOBASE PAIR UPSTREAM.
AUTHOR(S): ELION E A; WARNER J R
CORPORATE SOURCE: DEP. BIOCHEM., ALBERT EINSTEIN COLL. MED., BRONX, N.Y.
10461.
SOURCE: CELL, (1984 (RECD 1985)) 39 (3 PART 2), 663-674.
CODEN: CELLB5. ISSN: 0092-8674.
FILE SEGMENT: BA; OLD
LANGUAGE: English

L6 ANSWER 12 OF 13 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
TI REPLICON SIZE OF **YEAST** RIBOSOMAL DNA.
AB The ribosomal RNA of the **yeast** Saccharomyces cerevisiae are transcribed from a 9kbp [kilobase pair] stretch of DNA which is **reiterated** .apprx. 120-fold in a continuous array, .apprx. 360 .mu.m long, on chromosome XII. Although ARS activity has been detected in the repeat unit, the size and disposition of replicons along this array of identical genes has not been determined. Immobilized rRNA was used as a probe to examine the size of radioactively labeled rDNA replicons resolved on alkaline sucrose gradients. The replicons were uniformly sized, .apprx.

5 repeat units in length, and groups of 4 adjacent replicons may be activated simultaneously. Replicon initiation events are not determined solely by the recognition of specific DNA sequences that function as origins of replication.

ACCESSION NUMBER: 1984:347097 BIOSIS
DOCUMENT NUMBER: BA78:83577
TITLE: REPLICON SIZE OF **YEAST RIBOSOMAL DNA**.
AUTHOR(S): WALMSLEY R M; JOHNSTON L H; WILLIAMSON D H; OLIVER S G
CORPORATE SOURCE: DEP. BIOCHEM. AND APPLIED MOLECULAR BIOL., UNIV. MANCHESTER
INST. SCIENCE AND TECHNOL., P.O. BOX 88, MANCHESTER M60 1QD, U.K.
SOURCE: MOL GEN GENET, (1984) 195 (1-2), 260-266.
CODEN: MGGEAE. ISSN: 0026-8925.
FILE SEGMENT: BA; OLD
LANGUAGE: English

L6 ANSWER 13 OF 13 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

TI SIMPLE MENDELIAN INHERITANCE OF THE **REITERATED RIBOSOMAL DNA OF YEAST**.

AB A diploid strain of **yeast** (*Saccharomyces cerevisiae*) was heterozygous for 2 forms of the highly repetitious **ribosomal DNA**. These forms could be distinguished by the pattern of fragments produced after digestion with the site-specific restriction endonuclease EcoRI. The mode of inheritance of **ribosomal DNA** was determined by tetrad analysis. Of 14 tetrads analyzed, 12 clearly showed the **ribosomal DNA** forms segregating as a single Mendelian unit. The simplest interpretation of this result is that all of the approximately 100 copies of the **ribosomal DNA** genes of the **yeast** cell are located on 1 chromosome and that meiotic recombination within these genes is suppressed. Two of the 14 tetrads showed the segregation patterns expected as the result of mitotic recombination within the **ribosomal DNA**. [The DNA probe was prepared from an *Escherichia coli* strain].

ACCESSION NUMBER: 1978:158913 BIOSIS
DOCUMENT NUMBER: BA65:45913
TITLE: SIMPLE MENDELIAN INHERITANCE OF THE **REITERATED RIBOSOMAL DNA OF YEAST**.
AUTHOR(S): PETES T D; BOTSTEIN D
CORPORATE SOURCE: DEP. MICROBIOL., 920 E. 58TH ST., UNIV. CHIC., CHICAGO, ILL. 60637, USA.
SOURCE: PROC NATL ACAD SCI U S A, (1977) 74 (11), 5091-5095.
CODEN: PNASA6. ISSN: 0027-8424.
FILE SEGMENT: BA; OLD
LANGUAGE: English

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| NEWS | 36 | Dec 17 | TOXCENTER enhanced with additional content |
| NEWS | 37 | Dec 17 | Adis Clinical Trials Insight now available on STN |
| NEWS | 38 | Dec 30 | ISMEC no longer available |
| NEWS | 39 | Jan 13 | Indexing added to some pre-1967 records in CA/CAPLUS |
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| NEWS | 41 | Jan 21 | PHARMAML offering one free connect hour in February 2003 |
| NEWS | 42 | Jan 29 | Simultaneous left and right truncation added to COMPENDEX, ENERGY, INSPEC |
| NEWS | 43 | Feb 13 | CANCERLIT is no longer being updated |
| NEWS | 44 | Feb 24 | METADEX enhancements |
| NEWS | 45 | Feb 24 | PCTGEN now available on STN |

NEWS 46 Feb 24 TEMA now available on STN
 NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation
 NEWS 48 Feb 26 PCTFULL now contains images
 NEWS 49 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results

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=> s yeast and fermentation
 L1 40441 YEAST AND FERMENTATION

=> s l1 and xylose
L2 4099 L1 AND XYLOSE

=> s l2 and integrate yeast chromosome
L3 0 L2 AND INTEGRATE YEAST CHROMOSOME

=> s integrate yeast chromosome
L4 1 INTEGRATE YEAST CHROMOSOME

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TI Yeast derived vector contg. gene for antibiotic resistance - controlled by yeast or synthetic promoter, able to integrate with yeast chromosome.

AN 1985-304934 [49] WPIDS

CR 1986-332093 [50]; 1996-189959 [20]

AB EP 163491 A UPAB: 19960529

Vector includes a gene for resistance to an antibiotic normally able to kill a host yeast cell, and the gene is transcribed from a yeast or synthetic promoter sequence. The vector can be integrated into a chromosome of the yeast host.

The vector may also contain (1) a gene heterologous to the host and (2) a homologous sequence of the chromosome, inserted in such a way that no interference with host metabolism occurs.

USE/ADVANTAGE - Yeast cells transformed with the vectors express e.g. glucoamylase (able to convert starch to glucose which is then converted to CO₂ or EtOH, for use in dough making or brewing). Those expressing malate permease are useful in wine making because they can eliminate malic acid. The heterologous gene can also express a therapeutically useful protein, e.g. interferon. These vectors are stable over many generations even in the absence of selection.

Dwg.0/4

Dwg.0/4

ABEQ EP 163491 B UPAB: 19960428

A yeast cell transformed by integration into a chromosome thereof of vector DNA; characterised in that the host yeast cell is an industrial non-haploid yeast cell; in that the vector DNA comprises a gene for resistance to an antibiotic otherwise capable of killing said yeast cell, said gene being transcribed from a promoter sequence which is capable of promoting the expression of said antibiotic resistance gene at a level which confers antibiotic resistance to said cell; in that said vector DNA comprises a sequence homologous with a sequence of said chromosome and is integrated therein; and in that said vector DNA further comprises a gene for a desired heterologous protein.

Dwg.0/4

ACCESSION NUMBER: 1985-304934 [49] WPIDS

CROSS REFERENCE: 1986-332093 [50]; 1996-189959 [20]

DOC. NO. CPI: C1985-131759

TITLE: Yeast derived vector contg. gene for antibiotic resistance - controlled by yeast or synthetic promoter, able to integrate with yeast chromosome.

DERWENT CLASS: B04 D16

INVENTOR(S): YOCUM, R R

PATENT ASSIGNEE(S): (YOCU-I) YOCUM R R; (OMNI-N) OMNIGENE INC; (BIOY) BIOTECHNICA INT INC

COUNTRY COUNT: 7

PATENT INFORMATION:

| PATENT NO | KIND | DATE | WEEK | LA | PG |
|------------|------|----------|------------|----|----|
| EP 163491 | A | 19851204 | (198549) * | EN | 27 |
| AU 8542709 | A | 19851128 | (198604) | | |
| BR 8502400 | A | 19860121 | (198610) | | |
| FI 8502024 | A | 19851123 | (198611) | | |

JP 61040793 A 19860227 (198615)
 DK 8502241 A 19851123 (198617)
 EP 163491 B1 19960327 (199617) EN 20
 DE 3588096 G 19960502 (199623)
 CA 1338857 C 19970121 (199715)

APPLICATION DETAILS:

| PATENT NO | KIND | APPLICATION | DATE |
|------------|------|-----------------|----------|
| EP 163491 | B1 | EP 1985-303625 | 19850522 |
| DE 3588096 | G | DE 1985-3588096 | 19850522 |
| | | EP 1985-303625 | 19850522 |
| CA 1338857 | C | CA 1985-481908 | 19850521 |

FILING DETAILS:

| PATENT NO | KIND | PATENT NO |
|------------|------------|-----------|
| DE 3588096 | G Based on | EP 163491 |

PRIORITY APPLN. INFO: US 1984-612796 19840522